US ERA ARCHIVE DOCUMENT

DATE OUT: 14 Sept 2005

SUBJECT: EP [x] MP [] PRODUCT CHEMISTRY REVIEW

DP BARCODE No.: D318499

REG./File Symbol No.: 81598-R

PRODUCT NAME: Rotam Abamectin Technical

COMPANY: Rotam Ltd.

FOOD USE: [] PC CODE: 122804

Decision No. 350156 Integrated Formulation [x]

TO: RM #07, John Herbert/Thomas Harris

Insecticide-Rodenticide Branch Registration Division (7505C)

FROM: Bruce F. Kitchens, Chemist

Technical Review Branch

Registration Division (7505C)

INTRODUCTION:

The registrant, Rotam Limited, is responding to a previous product chemistry review for the proposed manufacturing use product, Rotam Abamectin Technical. In that review, it was determined that a revised Confidential Statement of Formula (CSF) would have to be submitted to correct an error. In addition, the registrant has submitted additional preliminary analysis data for Rotam Abamectin Technical. TRB has also been asked to determine if the proposed product is substantially similar to EPA Reg. No. 100-895 Abamectin Technical. The active ingredient in this product is Abamectin at a label nominal concentration of 98.0%. This product is intended for use in the manufacture of insecticide end use products. In support of this request, the registrant has submitted a revised basic Confidential Statement of Formula (CSF) dated 02 June 2005, a draft label, and product chemistry data contained in MRID#s 465557-01 and 465634-01. The data contained in MRID# 465634-01 is intended to replace data contained in MRID# 465503-01. The Technical Review Branch (TRB) has been asked to review this submission.

SUMMARY OF FINDINGS:

TRB has reviewed this submission and reports the following findings:

1. This product is produced from an integrated formulation process. This means that the active ingredient is the result of intended chemical reactions. In addition, the manufacturing process has been modified to produce the



higher purity product. See confidential appendix for details.

- 2. The impurity profile has been adequately described. The registrant has not declared any impurities of toxicological concern.
- 3. The nominal concentration of the active ingredient listed on the proposed basic CSF and the draft label are the same.
- 4. The active ingredients' certified limits as proposed on the basic CSF are acceptable.
- 5. The comparison of the cited and proposed product reveals that both products have the same active ingredient, both products have different active ingredient concentrations 90% vs 98%, and both products have four common impurities while the cited product has eight impurities not found in the proposed product. Wider certified limits for the cited product were proposed and accepted by the Agency. The justification for the wider certified limits was based on manufacturing process variability.

CONCLUSIONS:

TRB has reviewed this submission and concludes the following:

- 1. The basic formula CSF for the proposed manufacturing use product, Rotam Abamectin Technical dated 02 June 2005 is acceptable.
- 2. This submission satisfies the data requirements as specified in 40 CFR 158.155, 158.160, 158.162, 158.167, 158.170, 158.175, and 158.180 with respect to product identity and composition, description of materials used to produce the product, description of production process, discussion of formation of impurities, preliminary analysis, certified limits, and enforcement analytical method.
- 3. This product has been determined to be substantially similar to EPA Reg. No. 100-895 Abamectin Technical MK-936 from a product chemistry standpoint only. While the active ingredient nominal concentrations are different, the proposed product's active ingredient concentration falls within the active ingredient certified limits of the cited product.



PRODUCT CHEMISTRY DATA (GROUP A)

H	ical IDs/Manufacture/ ytical Information	Data Required Fulfilled	MRID No.
830-1550	Product Identity and Composition	Y	465634-01
830-1600	Description of Materials Used to Produce the Product	Y	465634-01
830-1620	Description of Production Process	Y	465634-01
830-1650	Description of Formulation Process	NA	
830-1670	Discussion of Impurities	Y	465634-01
830-1700	Preliminary Analysis	Y	465557-01
830-1750	Certified Limits	Y	465634-01
830-1800	Enforcement Analytical Method	Y	465634-01

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Pages	through are not included.
The ma	terial not included contains the following type o
Ide	entity of product inert ingredients.
Ide	entity of product impurities.
V Des	scription of the product manufacturing process.
Des	scription of quality control procedures.
Ide	entity of the source of product ingredients.
Sa:	les or other commercial/financial information.
A c	draft product label.
The	product confidential statement of formula.
In:	formation about a pending registration action.
FI	FRA registration data.
The	document is a duplicate of page(s)
	document is not responsive to the request.